



Food and Drug Administration  
10903 New Hampshire Avenue  
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Silver Spring, MD 20993-0002

April 29, 2015

Andon Health Co., Ltd  
Mr. Liu Yi  
President  
No 3, Jinping Street Ya An Road, Nankai District,  
Tianjin, P.R. China

Re: K141984

Trade/Device Name: Fully Automatic Electronic Blood Pressure Monitors Models KD-927  
and KD-928  
Regulation Number: 21 CFR 870.1130  
Regulation Name: Noninvasive blood pressure measurement system  
Regulatory Class: Class II  
Product Code: DXN  
Dated: March 13, 2015  
Received: March 16, 2015

Dear Mr. Liu Yi:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21

CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

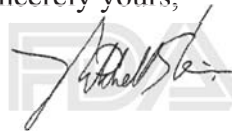
<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Bram D. Zuckerman", is written over a faint, large, light-gray watermark of the FDA seal.

for Bram D. Zuckerman, M.D.

Director

Division of Cardiovascular Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

## Statement of Indications for Use

**510(k) Number :** K141984

**Device name:** KD-927,KD-928 Fully Automatic Electronic Blood Pressure Monitor

**Indications for use:**

KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor are for use by medical professionals or at home and are non-invasive blood pressure measurement system intended to measure the diastolic and systolic blood pressures and pulse rate of an adult individual by using a non-invasive technique in which an inflatable cuff is wrapped around the upper arm. The cuff circumference is limited to 22cm-48cm.

Prescription use \_\_\_\_\_ AND/OR Over-The-Counter Use YES  
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

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## 510(k) Summary

510 (k) number: K141984

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirement of SMDA and 21 CFR 807.92.

### **1.0 submitter's information**

Name: Andon Health Co., Ltd.  
Address: No 3, Jinping Street Ya An Road, Nankai District, Tianjin,  
P.R. China  
Phone number: 86-22-6052 6161  
Fax number: 86-22-6052 6162  
Contact: Liu Yi  
Date of Application: 7/7/2014

### **2.0 Device information**

Trade name: KD-927,KD-928 Fully Automatic Electronic Blood Pressure  
Monitor  
Device name: KD-927,KD-928 Fully Automatic Electronic Blood Pressure  
Monitor  
Classification name: Noninvasive blood pressure measurement system

### **3.0 Classification**

Production code: DXN- Noninvasive blood pressure measurement system.  
Regulation number: 870.1130  
Classification: II  
Panel: Cardiovascular

### **4.0 Predicate device information**

Manufacturer: Andon Health Co., Ltd.  
Device: KD-936 Fully Automatic Wireless Blood Pressure Monitor  
510(k) number: K120672

## **5.0 Intended use**

KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor are for use by medical professionals or at home and are non-invasive blood pressure measurement system intended to measure the diastolic and systolic blood pressures and pulse rate of an adult individual by using a non-invasive technique in which an inflatable cuff is wrapped around the upper arm. The cuff circumference is limited to 22cm-48cm.

The intended use and the indication for use of KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor, as described in its labeling are the same as the predicate device KD-936(K120672) .

## **6.0 Device description**

KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor are designed and manufactured according to IEC80601-2-30--manual, electronic or automated sphygmomanometers.

The operational principle is based on oscillometric and silicon integrates pressure sensor technology. It can calculate the systolic and diastolic blood pressure, the measurements results can also be classified by the function of blood pressure classification indicator. If any irregular heartbeat is detected, it can be shown to the user. More over, it also obtains the function of averaging the measurement results. Achieves its function by an LCD or cell phone.

## **7.0 Summary comparing technological characteristics with predicate device**

<b>Technological Characteristics</b>	<b>Comparison result</b>
Design principle	Identical
Appearance	Similar
Patients contact Materials	Identical
Performance	Similar
Biocompatibility	Identical
Mechanical safety	Identical
Energy source	Similar
Standards met	Identical
Electrical safety	Identical
EMC	Identical
Function	Similar

## **8.0 Discussion of non-clinical and clinical test performed**

### **Non-clinical Tests have been done as follows:**

- a. Electromagnetic compatibility test according to IEC 60601-1-2;
- b. Electrical safety according test to IEC 60601-1 and IEC 60601-1-1
- c. Safety and performance characteristics of the test according to IEC 80601-2-30

None of the test demonstrates that KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor bring new questions of safety and effectiveness.

## **9.0 Performance summary**

Fully Automatic Electronic Blood Pressure Monitor conforms to the following standards:

- IEC 60601-1:2005/EN 60601-1:2006/AC:2010 (Medical electrical equipment -- Part 1: General requirements for basic safety and essential performance)
- IEC60601-1-2:2007/EN 60601-1-2:2007 /AC:2010 (Medical electrical equipment -- Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests)
- IEC 80601-2-30:2009+Cor.2010/EN 80601-2-30:2010(Medical electrical equipment –Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers)

## **10.0 Comparison to the predicate device and the conclusion**

Our device KD-927 and KD-928 Fully Automatic Electronic Blood Pressure Monitor are substantially equivalent to the Fully Automatic Wireless Electronic Blood Pressure Monitor KD-936 whose 510(k) number is K120672.

The KD-927 and KD-928 are very similar with the predicate device KD-936(K120672) in the intended use, the design principle, the material, the performance and the applicable standards, only their appearance, the memory time, the display components, electrical power, measure process, average measure function and Blood pressure non - rest condition index (BPNRI) function are different from the KD-936.

The conclusions drawn from the non-clinical and clinical tests demonstrate that the device is as safe, as effective, and performs as well as or better than the predicate device.